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## **Claims**

1	1.	A method for treating neurotrauma, said method comprising
2	administering	to a subject having neurotrauma a therapeutically effective amount
3	of a non-stere	oidal, anti-inflammatory drug (NSAID), analog, substituted form,
4	derivative, or	a pharmaceutically acceptable salt, ester, amide, or prodrug thereof.
l	2.	A method according to claim 1, wherein the NSAID is intrathecally
2	delivered.	
l	3.	A method according to claim 1, wherein the NSAID is
2	intraventricul	arly delivered.
l	4.	A method according to claim 1, wherein the non-steroidal anti-
2	inflammatory	drug comprises choline magnesium trisalicylate. 2189 511169
l	5.	A method according to claim 1, wherein the non-steroidal anti-
2	inflammatory	A method according to claim 1, wherein the non-steroidal anti- drug comprises sodium salicylate. 8515
		Aspir
1	6.	A method according to claim 1, wherein the non-steroidal anti-
2	inflammatory	drug comprises salicylamide. 8187 GHILL
		863 5' \
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		drug comprises salicylamide. 8187 SHIGH 514/165  863 514/165  RSPNICALLA 8190  SALICY SALICY  SALICY
		yd'. V.

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thereof.



1	7. A method according to claim 1, wherein the non-steroidal anti-
2	inflammatory drug comprises a deacetylated aspirin.
1	8. A method of treating diffuse axonal injury, said method comprising
2	administering to a subject having diffuse axonal injury a therapeutically effective
3	amount of a naturally occurring omega conotoxin, a functional fragment thereof, a
4	pharmacologically acceptable salt, ester, amide, or prodrug thereof.
1	9. A method according to claim 8, wherein the naturally occurring
2	omega conotoxin, functional fragment thereof, a pharmacologically acceptable
3	salt, ester, amide, or prodrug thereof is an N-type calcium channel blocker.
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1	10. A method according to claim 8, wherein the naturally occurring
2	omega conotoxin is selected from the group consisting essentially of GVIA and
3	MVII.
1	11. A method according to claim 8, wherein said administering step
2	further comprises intrathecally delivering the omega conotoxin, functional

fragment thereof, a pharmacologically acceptable salt, ester, amide, or prodrug

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caused by traumatic brain injury.



1	12. A method according to claim 8, wherein said administering step
2	further comprises intraventricularly delivering the omega conotoxin, functional
3	fragment thereof, a pharmacologically acceptable salt, ester, amide, or prodrug
4	thereof.
1	13. A method according to claim 8, wherein said administering step
2	further comprises delivering the omega conotoxin, functional fragment thereof, a
3	pharmacologically acceptable salt, ester, amide, or prodrug thereof to the subject
4	through an implantable pump.
1	14. A method according to claim 8, wherein said administering step
2	further comprises delivering the omega conotoxin, functional fragment thereof, a
3	pharmacologically acceptable salt, ester, amide, or prodrug thereof to the subject
4	through a spinal catheter.
1	15. A method according to claim 8, wherein the diffuse axonal injury is
2	a spastic disorder.

A method according to claim 15, wherein the spastic disorder is

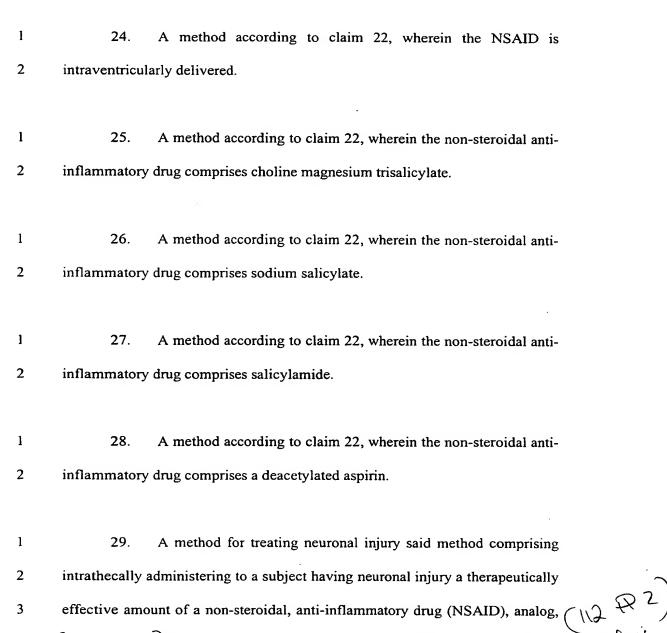
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1	17. A method according to claim 8 further including the step of
2	administering a non-steroidal anti-inflammatory drug to the subject.

- A method according to claim 17, wherein the non-steroidal anti-1 18. 2 inflammatory drug comprises choline magnesium trisalicylate.
- 1 19. A method according to claim 17, wherein the non-steroidal anti-2 inflammatory drug comprises sodium salicylate.
- 1 20. A method according to claim 17, wherein the non-steroidal anti-2 inflammatory drug comprises salicylamide.
- 1 21. A method according to claim 17, wherein the non-steroidal anti-2 inflammatory drug comprises a deacetylated aspirin.
- 1 22. A method for treating pain, said method comprising administering 2 to a subject having pain a therapeutically effective amount of a non-steroidal, anti-3 inflammatory drug (NSAID), analog, substituted form, derivative, or a 4 pharmaceutically acceptable salt, ester, amide, or prodrug thereof.
- 1 23. A method according to claim 22, wherein the NSAID is 2 intrathecally delivered.

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prodrug thereof.



substituted form derivative, or a pharmaceutically acceptable salt, ester, amide, or



1	30. A method according to claim 29, wherein the NSAID is
2	intraventricularly delivered.
1	31. A method according to claim 29, wherein the non-steroidal anti-
2	inflammatory drug comprises choline magnesium trisalicylate.
1	32. A method according to claim 29, wherein the non-steroidal anti-
2	inflammatory drug comprises sodium salicylate.
1	33. A method according to claim 29, wherein the non-steroidal anti-
2	inflammatory drug comprises salicylamide.
1	34. A method according to claim 29, wherein the non-steroidal anti-
2	inflammatory drug comprises a deacetylated aspirin.
1	35. A method according to claim 29, wherein the neuronal injury is
2	caused by Lupus, inflammatory neuropathy, infection, acquired disorders,

transverse myelitis, Parkinson's disease, CNS vasculitis, or Alzheimer's disease.